UV-sensitive syndrome

UV-sensitive syndrome is a condition that is characterized by sensitivity to the ultraviolet (UV) rays in sunlight. Even a small amount of sun exposure can cause a sunburn in affected individuals. In addition, these individuals can have freckles, dryness, or changes in coloring (pigmentation) on sun-exposed areas of skin after repeated exposure. Some people with UV-sensitive syndrome have small clusters of enlarged blood vessels just under the skin (telangiectasia), usually on the cheeks and nose. Although UV exposure can cause skin cancers, people with UV-sensitive syndrome do not have an increased risk of developing these forms of cancer compared with the general population.

Frequency

UV-sensitive syndrome appears to be a rare condition; only a small number of affected individuals have been reported in the scientific literature. However, this condition may be underdiagnosed.

Genetic Changes

UV-sensitive syndrome can result from mutations in the *ERCC6* gene (also known as the *CSB* gene), the *ERCC8* gene (also known as the *CSA* gene), or the *UVSSA* gene. These genes provide instructions for making proteins that are involved in repairing damaged DNA. DNA can be damaged by UV rays from the sun and by toxic chemicals, radiation, and unstable molecules called free radicals. Cells are usually able to fix DNA damage before it causes problems. If left uncorrected, DNA damage accumulates, which causes cells to malfunction and can lead to cell death.

Cells have several mechanisms to correct DNA damage. The CSB, CSA, and UVSSA proteins are involved in one mechanism that repairs damaged DNA within active genes (those genes undergoing gene transcription, the first step in protein production). When DNA in active genes is damaged, the enzyme that carries out gene transcription (RNA polymerase) gets stuck, and the process stalls. Researchers think that the CSB, CSA, and UVSSA proteins help remove RNA polymerase from the damaged site, so the DNA can be repaired.

Mutations in the *ERCC6*, *ERCC8*, or *UVSSA* genes lead to the production of an abnormal protein or the loss of the protein. If any of these proteins is not functioning normally, skin cells cannot repair DNA damage caused by UV rays, and transcription of damaged genes is blocked. However, it is unclear exactly how abnormalities in these proteins cause the signs and symptoms of UV-sensitive syndrome.

Inheritance Pattern

This condition is inherited in an autosomal recessive pattern, which means both copies of the gene in each cell have mutations. The parents of an individual with an autosomal recessive condition each carry one copy of the mutated gene, but they typically do not show signs and symptoms of the condition.

Other Names for This Condition

- ultraviolet sensitive syndrome
- UVSS

Diagnosis & Management

Genetic Testing

- Genetic Testing Registry: UV-sensitive syndrome https://www.ncbi.nlm.nih.gov/gtr/conditions/C1833561/
- Genetic Testing Registry: UV-sensitive syndrome 2 https://www.ncbi.nlm.nih.gov/qtr/conditions/C3553298/
- Genetic Testing Registry: UV-sensitive syndrome 3 https://www.ncbi.nlm.nih.gov/gtr/conditions/C3553328/

Other Diagnosis and Management Resources

- Merck Manual Home Health Edition: Sunburn http://www.merckmanuals.com/home/skin-disorders/sunlight-and-skin-damage/ sunburn
- World Health Organization: Sun Protection http://www.who.int/uv/sun_protection/en/

General Information from MedlinePlus

- Diagnostic Tests
 https://medlineplus.gov/diagnostictests.html
- Drug Therapy https://medlineplus.gov/drugtherapy.html
- Genetic Counseling https://medlineplus.gov/geneticcounseling.html
- Palliative Care https://medlineplus.gov/palliativecare.html
- Surgery and Rehabilitation https://medlineplus.gov/surgeryandrehabilitation.html

Additional Information & Resources

MedlinePlus

- Health Topic: Skin Conditions https://medlineplus.gov/skinconditions.html
- Health Topic: Sun Exposure https://medlineplus.gov/sunexposure.html

Genetic and Rare Diseases Information Center

 UV sensitive syndrome https://rarediseases.info.nih.gov/diseases/10947/uv-sensitive-syndrome

Educational Resources

- Disease InfoSearch: UV-sensitive syndrome http://www.diseaseinfosearch.org/UV-sensitive+syndrome/9441
- Disease InfoSearch: UV-SENSITIVE SYNDROME 2 http://www.diseaseinfosearch.org/UV-SENSITIVE+SYNDROME+2/9442
- Disease InfoSearch: UV-SENSITIVE SYNDROME 3
 http://www.diseaseinfosearch.org/UV-SENSITIVE+SYNDROME+3/9443
- MalaCards: uv-sensitive syndrome http://www.malacards.org/card/uv_sensitive_syndrome
- Merck Manual Home Health Edition: Overview of Sunlight and Skin Damage http://www.merckmanuals.com/home/skin-disorders/sunlight-and-skin-damage/ overview-of-sunlight-and-skin-damage
- Orphanet: UV-sensitive syndrome http://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=178338
- World Health Organization: UV Radiation http://www.who.int/uv/faq/whatisuv/en/index2.html

Patient Support and Advocacy Resources

 Canadian Skin Patient Alliance http://www.canadianskin.ca/en/

Scientific Articles on PubMed

PubMed

https://www.ncbi.nlm.nih.gov/pubmed?term=%28%28uv-sensitive+syndrome %5BTIAB%5D%29+OR+%28UVSS%5BTIAB%5D%29%29+AND+english%5Bla %5D+AND+human%5Bmh%5D+AND+%22last+3600+days%22%5Bdp%5D

OMIM

- UV-SENSITIVE SYNDROME 1 http://omim.org/entry/600630
- UV-SENSITIVE SYNDROME 2 http://omim.org/entry/614621
- UV-SENSITIVE SYNDROME 3 http://omim.org/entry/614640

Sources for This Summary

- Horibata K, Iwamoto Y, Kuraoka I, Jaspers NG, Kurimasa A, Oshimura M, Ichihashi M, Tanaka K. Complete absence of Cockayne syndrome group B gene product gives rise to UV-sensitive syndrome but not Cockayne syndrome. Proc Natl Acad Sci U S A. 2004 Oct 26;101(43):15410-5. Epub 2004 Oct 14.
 - Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/15486090 Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC524447/
- Itoh T, Fujiwara Y, Ono T, Yamaizumi M. UVs syndrome, a new general category of photosensitive disorder with defective DNA repair, is distinct from xeroderma pigmentosum variant and rodent complementation group I. Am J Hum Genet. 1995 Jun;56(6):1267-76.
 Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/7539208
 Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC1801097/
- Itoh T, Yamaizumi M, Ichihashi M, Hiro-Oka M, Matsui T, Matsuno M, Ono T. Clinical characteristics of three patients with UVs syndrome, a photosensitive disorder with defective DNA repair. Br J Dermatol. 1996 Jun;134(6):1147-50.
 Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/8763445
- Nardo T, Oneda R, Spivak G, Vaz B, Mortier L, Thomas P, Orioli D, Laugel V, Stary A, Hanawalt PC, Sarasin A, Stefanini M. A UV-sensitive syndrome patient with a specific CSA mutation reveals separable roles for CSA in response to UV and oxidative DNA damage. Proc Natl Acad Sci U S A. 2009 Apr 14;106(15):6209-14. doi: 10.1073/pnas.0902113106. Epub 2009 Mar 27. Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/19329487
 Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC2667150/
- Spivak G, Itoh T, Matsunaga T, Nikaido O, Hanawalt P, Yamaizumi M. Ultraviolet-sensitive syndrome cells are defective in transcription-coupled repair of cyclobutane pyrimidine dimers. DNA Repair (Amst). 2002 Aug 6;1(8):629-43.
 Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/12509286
- Zhang X, Horibata K, Saijo M, Ishigami C, Ukai A, Kanno S, Tahara H, Neilan EG, Honma M, Nohmi T, Yasui A, Tanaka K. Mutations in UVSSA cause UV-sensitive syndrome and destabilize ERCC6 in transcription-coupled DNA repair. Nat Genet. 2012 May;44(5):593-7. doi: 10.1038/ng.2228.
 - Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/22466612

Reprinted from Genetics Home Reference:

https://ghr.nlm.nih.gov/condition/uv-sensitive-syndrome

Reviewed: July 2012

Published: March 21, 2017

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